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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
09/709,249	11/10/2000	Alan H. Lazarus	701826-050990	3430
7:	590 05/17/2002		_	
Nixon Peabody LLP			EXAMINER	
101 Federal Street Boston, MA 02110			EWOLDT, GI	ERALD R
			ART UNIT	PAPER NUMBER
			1644	7
			DATE MAILED: 05/17/2002	/

Please find below and/or attached an Office communication concerning this application or proceeding.

Office Action Summary

Application No. 09/709,249

Applicant(s)

Lazarus et al.

Examiner

G.R. Ewoldt

Art Unit **1644**



The MAILING DATE of this communication appears	on the cover sheet with the correspondence address
Period for Reply	
A SHORTENED STATUTORY PERIOD FOR REPLY IS SET	TO EXPIRE 3 MONTH(S) FROM
THE MAILING DATE OF THIS COMMUNICATION. - Extensions of time may be available under the provisions of 37 CFR 1.136 (a). In	no event, however, may a reply be timely filed after SIX (6) MONTHS from the
mailing date of this communication.	
 If the period for reply specified above is less than thirty (30) days, a reply within the lift NO period for reply is specified above, the maximum statutory period will apply to the second sec	and will expire SIX (6) MONTHS from the mailing date of this communication.
 Failure to reply within the set or extended period for reply will, by statute, cause the Any reply received by the Office later than three months after the mailing date of 	he application to become ABANDONED (35 U.S.C. § 133).
earned patent term adjustment. See 37 CFR 1.704(b).	
Status 1 M Paragrains to communication(s) filed on Mar 6, 20	202
1) Responsive to communication(s) filed on <u>Mar 6, 20</u>	
	tion is non-final.
closed in accordance with the practice under Ex pa	except for formal matters, prosecution as to the merits is erte Quayle, 1935 C.D. 11; 453 O.G. 213.
Disposition of Claims	
4) 💢 Claim(s) <u>1-5</u>	is/are pending in the application.
4a) Of the above, claim(s)	is/are withdrawn from consideration.
5) Claim(s)	•
6) 💢 Claim(s) <u>1-5</u>	
7) Claim(s)	is/are objected to.
8)	are subject to restriction and/or election requirement.
Application Papers	
9) 💢 The specification is objected to by the Examiner.	
10) The drawing(s) filed on is/ard	e a) \square accepted or b) \square objected to by the Examiner.
	drawing(s) be held in abeyance. See 37 CFR 1.85(a).
	is: a) \square approved b) \square disapproved by the Examiner.
If approved, corrected drawings are required in reply	·
12) The oath or declaration is objected to by the Exam	niner.
Priority under 35 U.S.C. §§ 119 and 120	
13) Acknowledgement is made of a claim for foreign p	
a) ☐ All b) ☐ Some* c) ☐ None of:	
1. \square Certified copies of the priority documents ha	ve been received.
2. \square Certified copies of the priority documents ha	ve been received in Application No
application from the International Bur	
*See the attached detailed Office action for a list of the	
14) 🗓 Acknowledgement is made of a claim for domestic	
a) The translation of the foreign language provision	
15) Acknowledgement is made of a claim for domestic	c priority under 35 U.S.C. §§ 120 and/or 121.
Attachment(s)	N□
1) Notice of References Cited (PTO-892)	4) Interview Summary (PTO-413) Paper No(s).
2) Notice of Draftsperson's Patent Drawing Review (PTO-948)	5) Notice of Informal Patent Application (PTO-152)
3) X Information Disclosure Statement(s) (PTO-1449) Paper No(s)4	6) L Other:

DETAILED ACTION

- 1. Applicant's election of the antibody species L368 in Paper No. 6 is acknowledged. Because applicant did not distinctly and specifically point out the supposed errors in the restriction requirement, the election has been treated as an election without traverse (MPEP § 818.03(a)).
- 2. Claims 1-5 are pending and being acted upon.
- 3. The specification is objected to for the following informalities, $\beta 2$ -microglobulin is misspelled in a number of different ways in the specification. See for example, B-2 microglobulin (page 4), B-2M (page 5), $\beta 2M$ (page 11), $_2M$ (page 14), etc.

Appropriate corrections are required.

- 4. Claims 1-4 are objected to because of the following informalities:
- A) in Claims 1, 3, and 4, the phrase "an HLA alloimmune response from said patient," would more properly be written, "an HLA alloimmune response in said patient."
- B) in Claim 2, the phrase "wherein said at least one monoclonal antibody is," would more properly be written, "wherein said at least one monoclonal antibody is selected from the group consisting of."
- C) in Claim 3, the phrase "transfusing with the presensitizes platelets of step a) to a patient," would more properly be written, "transfusing the presensitizes platelets of step a) into a patient."
- D) in Claim 4, the phrase "at least two transfusions from said patient," would more properly be written, "at least two transfusions into said patient."

Appropriate correction is required.

5. The following is a quotation of the first paragraph of 35 U.S.C. 112:

The specification shall contain a written description of the invention, and of the manner and process of making and using it, in such full, clear, concise, and exact terms as to enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to make and use the same and shall set forth the best mode contemplated by the inventor of carrying out his invention.

6. Claims 1, 2, and 5 are rejected under 35 U.S.C. 112, first paragraph, because the specification, while being enabling for,

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a method for inhibiting an HLA autoimmune response to platelet transfusion, does not reasonably provide enablement for,

a method for preventing an HLA autoimmune response to platelet transfusion or preventing refractoriness to subsequent transfusions.

The specification disclosure is insufficient to enable one skilled in the art to practice the invention as claimed without an undue amount of experimentation. Undue experimentation must be considered in light of factors including: the breadth of the claims, the nature of the invention, the state of the prior art, the level of one of ordinary skill in the art, the level of predictability of the art, the amount of direction provided by the inventor, the existence of working examples, and the quantity of experimentation needed to make or use the invention.

Regarding "preventing" an alloimmune response or "preventing" refractoriness to subsequent transfusions, "prevent" is defined as an absolute term. To prevent is to keep from happening or to stop (Webster's Ninth New Collegiate Dictionary, 1990). It is disclosed in the specification, however, that the method of the instant claims functions only to inhibit an See for example, the section at page 7, alloimmune response. Monoclonal HLA-A2 antibody-treated platelets inhibit alloantibody production, or the section at page 9, Monoclonal antibodies to HLA Class I, but not to platelet-specific antigens, inhibit the alloimmune response. Further, the data disclosed in Figure 3 discloses a reduction in the alloimmune response, but not an absolute prevention of said response. Thus, the use of the method of the instant claims for the prevention of an alloimmune response, or the prevention of refractoriness to subsequent transfusions, must be considered highly unpredictable. said unpredictability, significant working examples would be required. As the relevant working example (results disclosed in Figure 3) indicates that the claimed method inhibits, but does not prevent an alloimmune response, the claimed method must be considered to require undue experimentation as there would be no particular expectation of success in achieving an absolute prevention of an alloimmune response or prevention of refractoriness to subsequent transfusions.

In re Wands, 858 F.2d at 737, 8 USPQ2d at 1404 (Fed. Cir. 1988) indicates that the more unpredictable an area is, the more specific enablement is necessary in order to satisfy the statute. In view of the quantity of experimentation necessary and the lack

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of sufficient working examples, it would take undue trials and errors to practice the claimed invention.

- 7. The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:
 - (a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.
- 8. Claims 3 and 4 are rejected under 35 U.S.C. 103(a) as being unpatentable over Crow et al. (March 1999, IDS) in view of Damjanovich et al. (1995) and Shalit et al. (1985).

Crow et al. teaches a method for inhibiting HLA alloimmune response to platelet transfusion, said method comprising the step of presensitizing platelets with a polyclonal antibody against HLA, a portion thereof, or $\beta 2\text{-microglobulin},$ wherein said platelets if administered to a patient inhibit an HLA alloimmune response in said patient (see particularly page 921, Results, Pretreatment of platelets with an anti-HLA alloantibody reduces their immunogenicity and Figure 1). Further, the reference teaches the inhibition of an alloimmune response after at least two transfusions (see particularly page 923, column 1, 7th paragraph).

The reference differs from the claimed invention in that it does not teach the use of the anti- $\beta 2$ -microglobulin antibody L368 in the claimed method.

Damjanovich et al. teaches that L368 is an anti- β 2-microglobulin monoclonal antibody (see particularly page 1122, column 2, Monoclonal Antibodies).

Shalit et al. teaches that monoclonal antibodies are more specific, more sensitive, and more accurate than polyclonal antibodies (see particularly Abstract and page 878, column 2, first paragraph).

It would have been *prima facie* obvious to one of ordinary skill in the art at the time the invention was made to perform method for inhibiting HLA alloimmune response to platelet transfusion, said method comprising the step of presensitizing platelets with an antibody against HLA, a portion thereof, or $\beta 2$ -

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microglobulin, wherein said platelets if administered to a patient inhibit an HLA alloimmune response in said patient and the inhibition of an alloimmune response after at least two transfusions, as taught by Crow et al., substituting the L368 anti- $\beta 2$ -microglobulin antibody as taught by Damjanovich et al., for the polyclonal antibody taught by Crow et al. One of ordinary skill in the art at the time the invention was made would have been motivated to make said substitution because monoclonal antibodies are more specific, more sensitive, and more accurate than polyclonal antibodies, as taught by Shalit et al.

9. No claim is allowed.

10. Any inquiry concerning this communication or earlier communications from the examiner should be directed to Dr. Gerald Ewoldt whose telephone number is (703) 308-9805. The examiner can normally be reached Monday through Thursday from 7:30 am to 5:30 pm. A message may be left on the examiner's voice mail service. If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Christina Chan can be reached on (703) 308-3973. Any inquiry of a general nature or relating to the status of this application should be directed to the Technology Center 1600 receptionist whose telephone number is (703) 308-0196.

Papers related to this application may be submitted to Technology Center 1600 by facsimile transmission. Papers should be faxed to Technology Center 1600 via the PTO Fax Center located in Crystal Mall 1. The faxing of such papers must conform with the notice published in the Official Gazette, 1096 OG 30 (November 15, 1989). The CM1 Fax Center telephone number is (703) 305-3014.

G.R. Ewoldt, Ph.D.

Patent Examiner

Technology Center 1600

May 16, 2002